

References

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Reply

We agree that transient dyspnea is common after injection of doses of adenosine sufficient to produce change in sinus rate at rest. However, in comparison studies, we (1) have shown that the dosage required to terminate paroxysmal supraventricular tachycardia in the same patient is usually lower and thus the sensation of dyspnea is much less common.

The effects of adenosine on respiration are still incompletely understood. Some authors (2) have concluded that adenosine is a central respiratory depressant rather than a stimulant as proposed by Watt. In our experience, most patients with severe asthma are

receiving a methylxanthine. Because methylxanthines in therapeutic doses block the electrophysiologic effects of adenosine, we have not treated any such patients. Two of our patients had a history of mild asthma that did not require long-term prophylactic therapy; neither reported any unusual side effects from adenosine. Finally, the unusually rapid elimination of adenosine after injection is due to rapid cellular uptake by a mechanism blocked by dipyridamole. In our study we excluded patients receiving this agent because we anticipated that markedly lower dosage requirements would be observed. Because alternative forms of therapy are usually available, one might question whether adenosine should be the drug of choice in patients receiving dipyridamole.

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